

MS for Asian Myrmecology

**Defensive function of petiole spines in queens and workers of the formicine ant
Polyrhachis lamellidens (Hymenoptera: Formicidae) against an ant predator, the
Japanese treefrog *Hyla japonica***

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ABSTRACT

The defensive function of petiole spines in queens and workers of the formicine ant *Polyrhachis lamellidens* was investigated using the ant predating tree frog *Hyla japonica*. Ant workers have hook-like large spines on their petiole while the queen petiole has only small slightly curved spines. Intact workers of *P. lamellidens* are unpalatable while workers without spines and intact queens are palatable, indicating that the spines of workers provide an effective defense against the treefrogs. Caste differences of spine morphology therefore affect the defense potential of the ants.

INTRODUCTION

Ants are social insects that live together with nestmates, the majority of which are wingless workers. Such insects may form an easy prey for insectivorous animals, and therefore must have an efficient defense system (Redford 1987, Peeters & Ito 2015). An effective defense system might contribute to the current success of ants in the terrestrial ecosystem (Buschinger & Maschwitz 1984, Peeters & Ito 2015). However, investigations on the defense of ants by using ant predators have been rarely carried out to date, except for our studies using treefrogs and chicks (Ito et al. 2004, Taniguchi et al. 2005a).

The formicine genus *Polyrhachis* is one of the biggest ant genera (Wilson 1976), including 697 species (Bolton 2015). Many species of *Polyrhachis* are characterized by large spines on the petiole and/or alitrunk (Dorow 1995). Spines are one of the typical defense devices in animals and plants (e.g. Mikolajewski & Rolff 2004, Inbar & Lev-Yadum 2005, Hanley et al. 2007). Hook-like spines on the petiole of the workers in some subgenera of *Polyrhachis* seem to be a very powerful defense apparatus (Fig. 1). The function of the spines in *Polyrhachis* is supposed to protect them against vertebrate predators (Buschinger & Maschwitz 1984), however, no experimental evidence exists so far. Such remarkable spines may function as a visual signal against predators. As shown in aquatic firefly larvae, a conspicuous visual signal in conjunction with deterring substances functions as an effective

multimodal aposematic anti-predator defense (Fu et al. 2007). Interestingly, such large spines were found in workers only whereas the queen's petiole has only small spines (Kohout 2014).

We compared the defensive function of the spines of *P. lamellidens* Fr. Smith between queens and workers, by using the Japanese tree frog, *Hyla japonica* (Günther), which is a common generalist predator occurring in western Japan (Hirai & Matsui 2000), and which feeds on many species of ants (Taniguchi et al. 2005a,b, Ito et al. 2009). First, we compared the petiolar spines between queens and workers. The fine structure of the spine surface was also observed by SEM to check whether there are openings for injecting eventual chemical substances from the spine. Then, we investigated whether the frogs are able to feed on *P. lamellidens* workers and queens. To verify the defensive function of spines of workers, we offered ablated workers without spines to the frogs. Furthermore, to check whether spines of workers function as a visual signal for predators, we offered workers without spines to frogs that had previously experienced workers with spines.

MATERIALS AND METHODS

Ants

Polyrhachis lamellidens is an uncommon ant in western Japan. Colonies nest in large holes of standing trees in woodland. Recently this species was assigned as an endangered species by the Japan Ministry of the Environment (2014). The nuptial flight occurs in October to November. Mated dealated queens thereafter enter the nest of *Camponotus japonicus* or *C. obscuripes* (Kohribe 1963, Furukawa et al. 2013) or hibernate in dead wood, and enter the host ant nest the next spring. The founding queens show temporary social parasitism (Kohribe 1963): the *P. lamellidens* queen kills a host queen, and subsequently the parasite workers replace the host workers. Colony size is relatively large, reaching more than 10,000 workers.

Morphology of petiole spines

The petiole was removed from 10 workers and 10 queens, and its width and height were measured using Motic Images Plus 2.1 after digital photography. The Welch - test in the statistical Package R v3.0.2 (R Development Core Team 2013) was used for statistical analysis of average petiole size between queens and workers, because the test for homogeneity of variance can be omitted. A worker petiole for scanning microscopy was detached from the body, gold coated in a Bal-Tec Sputter Coater SCD 050 and examined in a JEOL JSM-6360 scanning electron microscope.

Behavior of frogs against ants

We collected in total 105 mature frogs (length of snout tip to cloacal opening > 3.00 cm) in and around rice fields in Miki-cho, Kagawa Prefecture, western Japan, from April to August. They were kept in the laboratory in glass containers. Mealworms were given as prey. Before the experiment, a frog was transferred into a small cylindrical container (\varnothing 120 mm x height 75 mm). The bottom of the container was covered with plaster to provide humidity. Each frog was subjected to food deprivation during three days prior to the experiment. A colony fragment of *P. lamellidens* including workers, alate queens and broods was kept in a large polypropylene container in the laboratory.

We offered an intact worker, a worker with petiole spines experimentally ablated (Fig. 1C), or a queen to a frog in the cylindrical container. The number of frogs examined in this experiment was 60 frogs for intact workers, 30 frogs for workers without spines, and 15 frogs for queens. All frogs were used for this experiment only once. Just before each experiment, the petiole spines of workers were cut off by clippers, while the wings of alate queens were removed with forceps. According to former reports on frog predation behavior by using

Treefrogs (Taniguchi et al., 2005a) and toads (Brower et al., 1960), frog behavior against prey animals can be divided into following three categories: ignore, attack but not feed on, and attack then feed on. In our study, the second category was further divided: reject the prey just after the frog touched it, or spit out after the ant was put in the mouth. For each frog, feeding behavior was recorded as follow. (1) The frog attacked or ignored the ant. If a frog

ignored the ant during 10 min, we stopped the experiment. (2) If the frog attacked the ant, we recorded whether the ant was put in the mouth or whether the ant was rejected just after the frog touched it. (3) For the frogs that took the ant in their mouth, we recorded whether they fed on the ant or whether they spit it out. Furthermore, to check whether frogs consider the spines as a visual signal, we offered a worker with or without spines to frogs (both $N = 10$), that had experienced an ant with spines and spit it out the day before. For this experiment, we recorded whether a frog attacks or not. When the frogs ignored or refused ants, we gave them a mealworm or a small cockroach to know whether they were hungry or not. Frog behavior was analyzed by contingency table test as in Brower et al. (1960) and Taniguchi et al. (2006). For comparisons of the behavior against each type of ants, pairwise comparison with fisher exact test in R v3.0.2 (R Development Core Team 2013) was used.

RESULTS

Morphology of spines

The morphological characteristics of the petiole are markedly different between workers and queens (Fig. 1). Workers have a pair of large hook-like spines whereas the queen's petiole has a pair of short slightly curved spines. Spine width of workers is slightly wider than that of queens ($t = -3.2$, $df = 13.9$, $P = 0.006$), but the height is remarkably different between the two castes ($t = -19.3$, $df = 15.2$, $P = 4.081e-12$). SEM observation of the hook of the worker's petiolar spine (Fig. 2) shows that there are many pores with a diameter around $1\ \mu\text{m}$ on its surface. The tip of the hook of the worker's spine has no such pores, and there were no major openings for emitting eventual chemical substances.

Behavior of frogs against ants

In total, 75 of 105 frogs that had no prior experience with ants in the laboratory attacked the ant (Table 1). The proportion of frogs that ignored ants was not statistically different among intact workers, mutilated workers, and queens. All 30 frogs that ignored ants

132 did feed on mealworm or cockroach, indicating that the frogs avoid the ants as prey. Among
133 the 75 frogs that attacked the ant, 15 frogs that attacked an intact worker and 8 frogs that
134 attacked an ablated worker refused the ant: they stopped attacking after their tongue touched
135 the ant. These frogs did feed on a mealworm or cockroach just after the experiment,
136 indicating that they were hungry but chemical substances or physical properties of the
137 cuticular surface including spines have some defensive function. The proportion of refusing
138 frogs was not statistically different with respect to their refusing of intact workers, ablated
139 workers, and intact queens. The remaining 52 frogs took ants in their mouth (Table 1). The
140 majority of frogs (27 out of 30 frogs) that put an intact worker into the mouth spit it out. When
141 the frogs took an intact worker, the spines often stuck in their mouth. Four frogs spent time
142 and effort to remove the ant from their mouth. All ants vomited by the frogs died. In
143 contrast, all but one frog fed on an ablated worker ($N = 14$). All frogs that took an intact
144 queen in their mouth ($N = 8$) ate it without vomiting. The feeding ratio of the intact workers
145 was significantly lower than that of both the ablated workers and queens (pairwise
146 comparison with fisher test, both $P < 0.00001$), indicating that the spines of *P. lamellidens*
147 workers do function as an effective defense against the predators.

148 Among 20 frogs that spit out the intact worker in the first experiment mentioned above
149 (Table 1), seven of 10 frogs who were offered the intact workers, and nine of 10 frogs who
150 were offered ablated workers ignored the ant (lower part of Table 1). The behavioral
151 response by frogs who experienced intact workers previously is significantly different from the
152 frogs before the experience (against intact workers, $P = 0.04$, against ablated workers, $P =$
153 0.0016), indicating that the frogs learned to recognize unpalatable prey. In this experiment,
154 both ablated ants and intact ants were ignored in a similar ratio, indicating that the spines
155 have no function as signals of bad prey for frogs.

157 DISCUSSION

158 The huge spines on the petiole of *P. lamellidens* workers have a strong defensive
159 effect against tree frogs, and that the frogs can learn to recognize unpalatable prey. To our

knowledge, this is the first experimental evidence on the defensive function of ant spines against predators. Defensive spines in insects sometimes can introduce toxins into another animals, e.g. larvae of Limacodidae (Rothschild et al. 1970, Murphy et al. 2010). SEM observation of the petiole indicated that there are many pores, which probably correspond with the opening of subepithelial glands. These glands are distributed over the whole body in several ant species (Gobin et al. 2003). It is unlikely that the tiny pores on the surface of the petiole spines of *P. lamellidens* emit toxic substances for defense, as they represent single cell openings, that cannot emit sufficiently large amounts of substance. Although we do not yet know the function of these glands, the defensive function of the spines seems based on their mere physical characteristics.

Spines of queens are small and have no defensive effect against the tree frogs. Huge spines seem to hinder for flying: generally ant queens have smaller spines if compared to conspecific workers (Kohout 2014, Peeters & Ito 2015). Furthermore, activity outside nests by queens is generally limited to just before and after the nuptial flight. Thus, such strong defensive apparatus against predators is not important in comparison to workers.

Our observations indicate that the tree frogs learn to recognize the unsuitable prey and avoid it as shown by Taniguchi *et al.* (2005a). However, the frogs that experienced the intact worker subsequently ignored not only intact workers but also the ablated workers (see lower part of Table 1), which indicates that the ant spines alone do not seem sufficient for the frogs to recognize unpalatable prey. How the tree frogs recognize such unpalatable prey is still unknown and will be studied in a future project.

The occurrence of such spines in *Polyrhachis* might contribute to the current prevalence of this genus. The defense function may be especially important for arboreal life where many vertebrate predators like tree lizards and birds forage. In contrast, workers of *Tetramorium tsushimae* Emery, *Crematogaster osakensis* Forel, and *Pheidole fervida* Fr. Smith which have small propodeal spines, are frequently eaten by tree frogs (Taniguchi et al. 2005b, Ito et al. 2009). At least against tree frogs, small spines are not functional as

defensive apparatus, however, it is possible that small spines are effective against the other predators.

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REFERENCES

- Bolton B, 2015. AntCat. An online catalog of the ants of the world. [Http://www.antcat.org](http://www.antcat.org).
- Brower LP, Brower JVZ and Westcott PW, 1960. Experimental studies of mimicry. 5. The reactions of toads (*Bufo terrestris*) to bumblebees (*Bombus americanorum*) and their robberfly mimics (*Mallophora bomboides*), with a discussion of aggressive mimicry. *The American Naturalist* 94:343-355.
- Buschinger A and Maschwitz U, 1984. Defensive behavior and defensive mechanisms in ants. In: *Defensive Mechanisms in Social Insects* (H.R. Hermann, Ed), Praeger Scientific, New York, pp 95-150
- Dorow WHO, 1995. Revision of the ant genus *Polyrhachis* Smith, 1857 (Hymenoptera: Formicidae: Formicinae) on subgenus level with keys, checklist of species and Bibliography. *Courier Forschungsinstitut Senckenberg* 185: 1-113.
- Furukawa K, Ohbe T and Ito F, 2012. Collection of a *Polyrhachis lamellidens* queen from a nest of *Camponotus obscuripes* (Hymenoptera: Formicidae). *Kagawa Seibutsu* 39: 61-62. In Japanese with English abstract.
- Gobin B, Ito F and Billen J, 2003. The subepithelial gland in ants: a novel exocrine gland closely associated with the cuticle surface. *Acta Zoologica* 84: 285-291

213 Hanley ME, Lamont BB, Fairbanks MM and Rafferty CM, 2007. Plant structural traits and their
 214 role in anti-herbivore defence. *Perspectives of Plant Ecology Evolution and Systematics*
 215 8: 157–178.

216 Hirai T and Matsui M, 2000. Feeding habits of the Japanese treefrog, *Hyla japonica*, in the
 217 reproductive season. *Zoologica Scripta* 17: 977-982

218 Hölldobler B and Wilson EO, 1990. *The Ants*. Harvard University Press, Cambridge, MA.
 219 732 pp

220 Inbar M and Lev-Yadun M, 2005. Conspicuous and aposematic spines in the animal
 221 kingdom. *Naturwissenschaften* 92: 170-172.

222 Ito F, Hashim R, Yek SH, Kaufmann E, Akino T and Billen J, 2004. Spectacular Batesian
 223 mimicry in ants. *Naturwissenschaften* 91: 481-484

224 Ito F, Okaue M and Ichikawa T, 2009. A note on prey composition of the Japanese treefrog,
 225 *Hyla japonica*, in an area invaded by Argentine ants, *Linepithema humile*, in Hiroshima
 226 Prefecture, western Japan (Hymenoptera: Formicidae). *Myrmecological News* 12:
 227 35-39.

228 Kohout R, 2014. A review of the subgenus *Polyrhachis* (*Polyrhachis*) Fr. Smith
 229 (Hymenoptera: Formicidae: Formicinae) with keys and description of a new species.
 230 *Asian Myrmecology* 6: 1-31.

231 Kohriba O, 1963. A parasitic life of *Polyrhachis lamellidens* F. Smith (Hymenoptera,
 232 Formicidae), first report. *Kontyu* 31: 200-209. In Japanese with English summary.

233 Mikolajewski DJ and Rolff J, 2004. Benefits of morphological defence demonstrated by
 234 direct manipulation in larval dragonflies. *Evolutionary Ecology Research* 6: 619-626

235 Murphy SM, Leahy SM, Williams LS and Lill JT, 2010. Stinging spines protect slug caterpillars
 236 (Limacodidae) from multiple generalist predators *Behavioral Ecology* 21: 153-160.

237 Peeters C and Ito F, 2015. Wingless and dwarf workers underlie the ecological success of
 238 ants (Hymenoptera, Formicidae). *Myrmecological News* 21: 117-130.

239 R Development Core Team, 2013. The statistical Package R v3.0.2.

240 Redford KH, 1987. Ants and termites as food. Patterns of mammalian myrmecophagy. In
241 Genoways HH eds. Current Mammalogy, pp 349-399, Springer
242 Rothschild M, Reichstein T, von Euw J, Alpin R and Harman RRM, 1970. Toxic Lepidoptera.
243 *Toxicon* 8: 293-296.
244 Taniguchi K, Maruyama M, Ichikawa T and Ito F, 2005a. A case of Batesian mimicry
245 between myrmecophilous staphylinid beetle, *Pella comes*, and its host ant, *Lasius*
246 (*Dendrolasius*) *spathepus*: an experiment using the Japanese treefrog *Hyla japonica* as
247 a real predator. *Insectes Sociaux* 52 : 320-322
248 Taniguchi K, Tanaka H and Ito F, 2005b. Ants as prey of the Japanese treefrog *Hyla japonica*
249 in an urban area of western Japan. *Sociobiology* 45: 495-502
250 Wilson EO, 1976. Which are the most prevalent ant genera? *Studia Entomologia* 19: 187-200.
251

Table 1. Behavioral responses of the Japanese tree frogs against queens and two types of *Polyrhachis lamellidens* workers. The different letters in each column refer to a significant difference (Ryan-test).

					from attacked ants:		from ants that were put in mouth:		
	Ignored	Attacked	Total	Refused	Put in mouth	Total	Fed on	Spit	Total
Frogs without experience:									
intact workers	15	45	60 ^a	15	30	45 ^a	3	27	30 ^a
mutilated workers	8	22	30 ^{a,c}	8	14	22 ^a	13	1	14 ^b
queens	7	8	15 ^{a,b}	0	8	8 ^a	8	0	8 ^b
Total	30	75	105	23	52	75	24	28	52
Frogs that spit out intact workers:									
intact workers	7	3	10 ^{b,c}						
mutilated workers	9	1	10 ^b						
Total	16	4	20						

Figure Legends

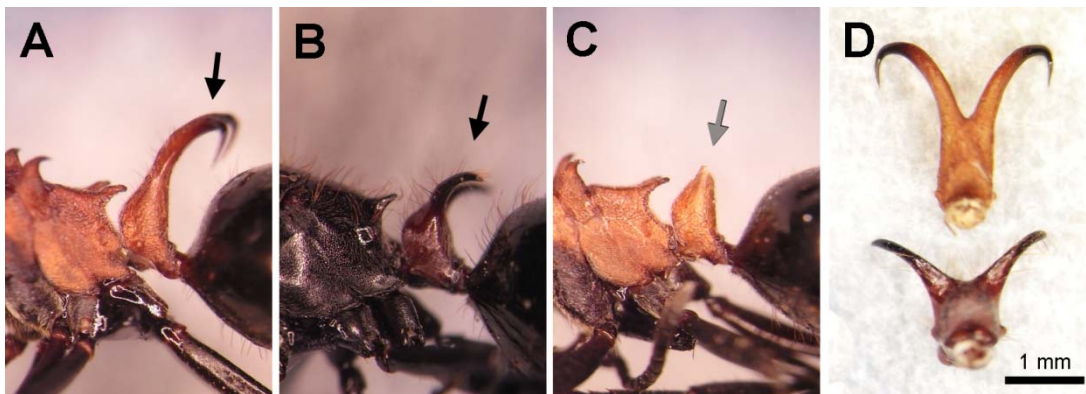


Fig. 1. Petiole profiles of queen (A), intact worker (B), and mutilated worker (C), and frontal view of petiole of queen (D top) and worker (D bottom) of *Polyrhachis lamellidens*. Black arrows indicate petiolar spine, grey arrow indicates mutilated spine of worker petiole.

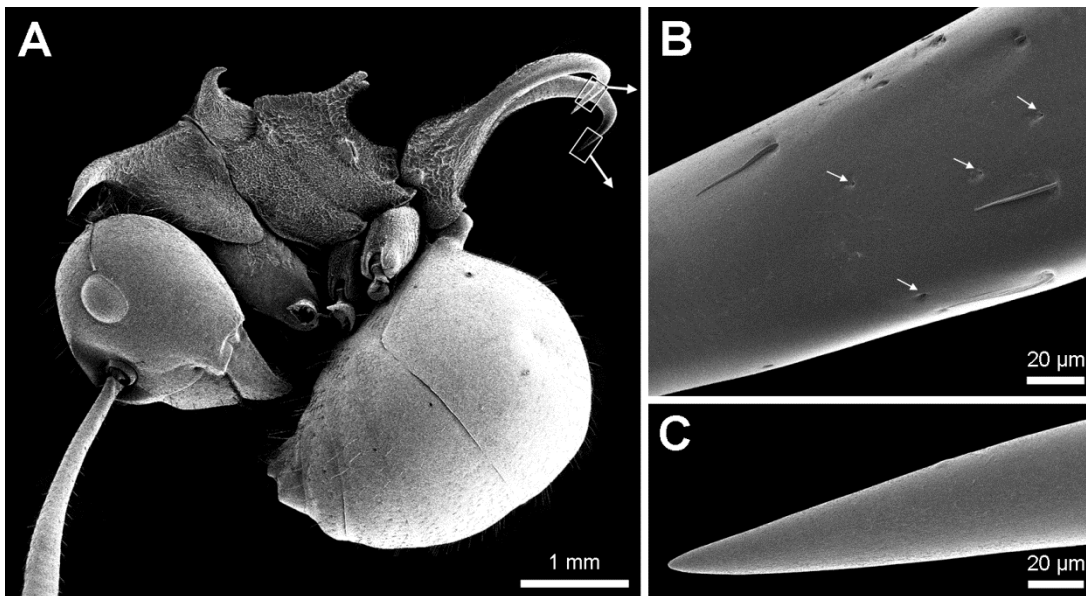


Fig. 2. SEM appearance of worker petiole in *Polyrhachis lamellidens*. Small arrows in B indicate pore openings of subepidermal gland ducts.